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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/031,342	03/26/2002	Torben Halkier	3631-0118P	6459
2292	7590	06/30/2004	EXAMINER	
BIRCH STEWART KOLASCH & BIRCH PO BOX 747 FALLS CHURCH, VA 22040-0747			BELYAVSKIY, MICHAEL A	
			ART UNIT	PAPER NUMBER
			1644	

DATE MAILED: 06/30/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary	Application No.	Applicant(s)	
	10/031,342	HALKIER ET AL.	
	Examiner	Art Unit	
	Michail A Belyavskyi	1644	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED. (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 14 August 2002.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1 and 53-101 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☐ Claim(s) _____ is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☒ Claim(s) 1 and 53-101 are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. _____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|---|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152) |
| Paper No(s)/Mail Date _____ | 6) <input type="checkbox"/> Other: _____ |

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DETAILED ACTION

1. Applicant's amendment, filed 08/14/02 is acknowledged.

Claims 1 and 53-101 are pending.

Restriction

2. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in response to this action, to elect a single invention to which the claims must be restricted:

I. Claim 1, drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising administering at least one GDF-8 polypeptide or subsequence thereof **and** at least one GDF-8 analogue.

II. Claim 1, drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising administering at least one GDF-8 polypeptide or subsequence thereof **or** at least one GDF-8 analogue.

III. Claims 53-74 and 80, drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising presentation to the animal's immune system at least one GDF-8 analogue, wherein a foreign T_H epitope is introduced in **SEQ ID NO:11**.

IV. Claims 53-74 and 80, drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising presentation to the animal's immune system at least one GDF-8 analogue, wherein a foreign T_H epitope is introduced in **SEQ ID NO:12**.

V. Claims 53, 75-79, drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising presentation to the animal's immune system at least one GDF-8 analogue, wherein a foreign T_H epitope is introduced in **SEQ ID NO:11**, wherein presentation of said analogue is effected by introducing nucleic acid encoding the GDF-8 analogue.

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VI. Claims 53, 75-79, drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising presentation to the animal's immune system at least one GDF-8 analogue, wherein a foreign T_H epitope is introduced in **SEQ ID NO:12**, wherein presentation of said analogue is effected by introducing nucleic acid encoding the GDF-8 analogue.

VII. Claims 81-83 drawn to a GDF analogue wherein a foreign T_H epitope is introduced in **SEQ ID NO:11** and an immunogenic composition comprising said analogue.

VIII. Claims 81-83 drawn to a GDF analogue wherein a foreign T_H epitope is introduced in **SEQ ID NO:12** and an immunogenic composition comprising said analogue.

IX. Claims 84-94, 99 ,100 and 101 drawn to a nucleic acid fragment which encodes a GDF-8 analogue wherein a foreign T_H epitope is introduced in **SEQ ID NO:11**; a vector carrying said nucleic acid fragment and a host cell transformed by said vector and a composition comprising said nucleic acid fragment.

X. Claims 84-94, 99 ,100 and 101 drawn to a nucleic acid fragment which encodes a GDF-8 analogue wherein a foreign T_H epitope is introduced in **SEQ ID NO:12**; a vector carrying said nucleic acid fragment and a host cell transformed by said vector and a composition comprising said nucleic acid fragment.

XI. Claims 53 and 95-98 drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising presentation to the animal's immune system at least one GDF-8 analogue, wherein a foreign T_H epitope is introduced in **SEQ ID NO:11**, wherein presentation of said analogue is effected by administering a non-pathogenic micro-organism or virus.

XII. Claims 53 and 95-98 drawn to a method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity comprising presentation to the animal's immune system at least one GDF-8 analogue, wherein a foreign T_H epitope is introduced in **SEQ ID NO:12**, wherein presentation of said analogue is effected by administering a non-pathogenic micro-organism or virus.

3. The inventions listed as Groups I-XLIX do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

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As was also found in the International Search Report, the Invention of Group I was found to have no special technical feature that defined the contribution over the prior art of WO'99/02667

WO'667 the method of using myostatin, also known as GDF-8 in order to down-regulate GDF-8 and thus increase muscle mass in a mammal by the production of antibodies against the GDF-8, by administering to the animal GDF-8 polypeptide or by an oligonucleotides.

Since Applicant's Inventions do not contribute a special technical feature when viewed over the prior art they do not have a single general inventive concept and so lack unity of invention.

Species Election

4. This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

5. If any one of the Groups III-XII is elected, applicant is required to elect a specific method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity wherein a T_H epitope is introduced into a specific position for example at position 18-41, as recited in claim 53.

These species are distinct because a specific method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity wherein a T_H epitope is introduced into a specific position for example at position 18-41, as recited in claim 53 differ with respect to the structure and mode of action of the specific GDF-8 analogue, thus each specific method employing a specific analogue represent a patentably distinct subject matter. The examination of species would require different searches in the scientific literature.

In addition:

6. If groups III or IV is elected, applicant is required to elect a specific method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity wherein: (i) specific T_H epitope is selected from the group recited in claim 62; (ii) first moiety is specific for one antigen as recited in claim 63; (iii) specific second moiety is selected from the group recited in claim 64 or 65; (iv) specific third moiety is selected from the group recited in claim 66; (v) specific GDF-8 analogue is derived from the group as recited in claim 67.

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These species are distinct because a specific method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity wherein (i) specific T_H epitope is selected from the group recited in claim 62; (ii) first moiety is specific for one antigen as recited in claim 63; (iii) specific second moiety is selected from the group recited in claim 64 or 65; (iv) specific third moiety is selected from the group recited in claim 66; (v) specific GDF-8 analogue is derived from the group as recited in claim 67 differ with respect to the structure and mode of action of the specific GDF-8 analogue, thus each specific method employing a specific analogue represent a patentably distinct subject matter. The examination of species would require different searches in the scientific literature.

7. If Groups V or VI is elected, applicant is required to elect a specific method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity wherein a specific type of nucleic acid is selected from the group recited in claim 75, for example naked DNA DNA formulated with charged lipids etc).

These species are distinct because a specific method for in vivo down regulation of growth differentiation factor 8 (GDF-8) activity wherein activity wherein a specific type of nucleic acid is selected from the group recited in claim 75, for example naked DNA DNA formulated with charged lipids etc) differ with respect to the structure of the nucleic acid sequences and composition thereof, thus each specific method employing a specific analogue represent a patentably distinct subject matter. The examination of species would require different searches in the scientific literature.

Applicant is required under 35 U.S.C. § 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable.

8. Applicant is advised that a response to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 C.F.R. § 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. M.P.E.P. § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the

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examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. § 103 of the other invention.

9. Applicant is advised that the response to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed.

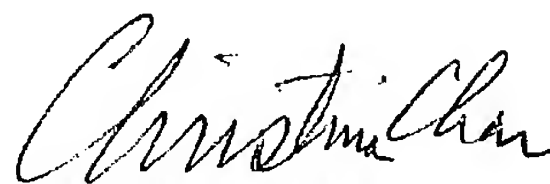
A telephone call was made to Leonard Svenson on 6/23/04 to request an oral election to the above restriction requirement, but did not result in an election being made.

10. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Michail Belyavskyi whose telephone number is 571/ 272-0840. The examiner can normally be reached Monday through Friday from 9:00 AM to 5:30 PM. A message may be left on the examiner's voice mail service. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christina Chan can be reached on 571/ 272-0841.

The fax number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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